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APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	CONFIRMATION NO.
10/667,133	09/19/2003	Howard M. Johnson	UF-243XD1	7182
23557	7590 03/31/2005		EXAMINER	
SALIWANCHIK LLOYD & SALIWANCHIK A PROFESSIONAL ASSOCIATION			GALVEZ, JAMES JASON	
PO BOX 14			ART UNIT	PAPER NUMBER
GAINESVI	LLE, FL 32614-2950	1647		
			DATE MAILED: 03/31/2005	5

Please find below and/or attached an Office communication concerning this application or proceeding.

		<i>\(\)</i>			
	Application No.	Applicant(s)			
Office Anti-e O	10/667,133	JOHNSON ET AL.			
Office Action Summary	Examiner	Art Unit			
	J. Jason Galvez	1647			
The MAILING DATE of this communicat Period for Reply	ion appears on the cover sheet w	th the correspondence address			
A SHORTENED STATUTORY PERIOD FOR THE MAILING DATE OF THIS COMMUNICA - Extensions of time may be available under the provisions of 37 after SIX (6) MONTHS from the mailing date of this communic - If the period for reply specified above is less than thirty (30) da - If NO period for reply is specified above, the maximum statuto. - Failure to reply within the set or extended period for reply will, Any reply received by the Office later than three months after the earned patent term adjustment. See 37 CFR 1.704(b).	TION. 7 CFR 1.136(a). In no event, however, may a ration. rys, a reply within the statutory minimum of thir y period will apply and will expire SIX (6) MON by statute, cause the application to become AB	eply be timely filed by (30) days will be considered timely. ITHS from the mailing date of this communication. SANDONED (35 U.S.C. § 133).			
Status					
1) Responsive to communication(s) filed o	n <u>9/30/2004</u> .				
	☐ This action is non-final.				
3) Since this application is in condition for	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is				
closed in accordance with the practice t	under <i>Ex parte Quayle</i> , 1935 C.D). 11, 453 O.G. 213.			
Disposition of Claims					
4) ⊠ Claim(s) 20-35 is/are pending in the appear 4a) Of the above claim(s) is/are versions 5) □ Claim(s) is/are allowed. 6) □ Claim(s) is/are rejected. 7) □ Claim(s) is/are objected to. 8) ⊠ Claim(s) 20-35 are subject to restriction	vithdrawn from consideration.				
Application Papers					
9)☐ The specification is objected to by the E	xaminer.				
10) The drawing(s) filed on is/are: a)	D)☐ The drawing(s) filed on is/are: a)☐ accepted or b)☐ objected to by the Examiner.				
Applicant may not request that any objection	-,,	• •			
Replacement drawing sheet(s) including the	•	• •			
11)☐ The oath or declaration is objected to by	the Examiner. Note the attached	Office Action of form P1O-152.			
Priority under 35 U.S.C. § 119					
12) Acknowledgment is made of a claim for a a) All b) Some * c) None of: 1. Certified copies of the priority doc 2. Certified copies of the priority doc 3. Copies of the certified copies of the application from the International * See the attached detailed Office action for	cuments have been received. cuments have been received in A he priority documents have been Bureau (PCT Rule 17.2(a)).	pplication No received in this National Stage			
Attachment(s)	_				
 Notice of References Cited (PTO-892) Notice of Draftsperson's Patent Drawing Review (PTO-93) Information Disclosure Statement(s) (PTO-1449 or PTO Paper No(s)/Mail Date 	948) Paper No(s	Summary (PTO-413) S)/Mail Date nformal Patent Application (PTO-152) 			

DETAILED ACTION

Election/Restrictions

Restriction to one of the following inventions is required under 35 U.S.C. 121:

- 1. Claims 20-35, drawn to a method of suppression or inhibition of allergenspecific IgE production by administering Interferon-tau, classified in class 514, subclass 2.
- Claims 20-35, drawn to a method of suppression or inhibition of allergenspecific IgE production by administering chimeric Interferon-tau, classified in class 514, subclass 2.

The inventions are distinct from one another with explanations as follows:

Inventions 1 and 2 are each unrelated to one another. Inventions are unrelated if it can be shown that they are not disclosed as capable of use together and they have different modes of operation, different functions, or different effects (MPEP § 806.04, MPEP § 808.01). In the instant case the different inventions are not disclosed as capable of use together. Furthermore, the inventions are directed to methods that use different molecules. The method invention 1 uses Interferon-tau whereas the method of invention 2 uses chimeric Interferon-tau. Chimeric Interferon-tau is structurally, and possibly functionally, different from unmodified Interferon-tau.

Because these inventions are distinct for the reasons given above and have acquired a separate status in the art as shown by their separate search requirement based on particular aspects of the inventions, *e.g.* invention 1 uses Interferon-tau and

invention 2 uses chimeric Interferon-tau, it would impose a serious burden on the Examiner and USPTO resources to search the inventions together.

This application contains claims directed to the following patentably distinct species of the claimed invention: routes of administration

- A) oral
- B) parenteral
- C) subcutaneous
- D) intravenous

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claim 1, <u>for example</u>, is generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

This application contains claims directed to the following patentably distinct species of the claimed invention: allergic conditions

- E) rhinitis
- F) atopic dermatitis
- G) bronchial asthma
- H) food allergy

Applicant is required under 35 U.S.C. 121 to elect a single disclosed species for prosecution on the merits to which the claims shall be restricted if no generic claim is finally held to be allowable. Currently, claim 23, for example, is generic.

Applicant is advised that a reply to this requirement must include an identification of the species that is elected consonant with this requirement, and a listing of all claims readable thereon, including any claims subsequently added. An argument that a claim is allowable or that all claims are generic is considered nonresponsive unless accompanied by an election.

Upon the allowance of a generic claim, applicant will be entitled to consideration of claims to additional species which are written in dependent form or otherwise include all the limitations of an allowed generic claim as provided by 37 CFR 1.141. If claims are added after the election, applicant must indicate which are readable upon the elected species. MPEP § 809.02(a).

Should applicant traverse on the ground that the species are not patentably distinct, applicant should submit evidence or identify such evidence now of record showing the species to be obvious variants or clearly admit on the record that this is the case. In either instance, if the examiner finds one of the inventions unpatentable over the prior art, the evidence or admission may be used in a rejection under 35 U.S.C. 103(a) of the other invention.

Applicant is advised that the reply to this requirement to be complete must include an election of the invention and an election of species to be examined even though the requirement may be traversed (37 CFR 1.143). Applicant is required to elect invention 1 or 2 and a species from A-D and E-H to be considered fully responsive.

Applicant is reminded that upon the cancellation of claims to a non-elected invention, the inventorship must be amended in compliance with 37 CFR 1.48(b) if one or more of the currently named inventors is no longer an inventor of at least one claim remaining in the application. Any amendment of inventorship must be accompanied by a request under 37 CFR 1.48(b) and by the fee required under 37 CFR 1.17(i).

Art Unit: 1647

Conclusion

Any inquiry concerning this communication or earlier communications from the examiner should be directed to **J. Jason Galvez, Ph.D.** whose telephone number is **571-272-2935**. The examiner can normally be reached Monday through Friday 9 AM to 5 PM. If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, **Brenda Brumback, Ph.D.** can be reached at **571-272-0887**.

The fax phone number for the organization where this application or proceeding is assigned is 571-273-8300. Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free).

JJG 3/30/2005

Bridget E. Bunner